## **Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

1. (currently amended) A <u>method of imaging or treating a tumor in a mammal</u> comprising:

administering an effective amount of a compound of the formula (I) to said mammal:

$$\begin{bmatrix} X & O \\ | & | \\ | Co | - [C - Y - Det] \end{bmatrix}$$
(I)

wherein the moiety is cobalamin, C is the residue of a monocarboxylic acid of cobalamin, X is CN, OH, methyl or adenosyl, Y is a linking group and Det is a detectable chelating group comprising a radionuclide or a paramagnetic metal ion; and optionally detecting the compound of formula I in a tumor of the mammal.

- 2. (currently amended) The <u>compound method</u> of claim 1 wherein the radionuclide is a metallic radioisotope.
- 3. (currently amended) The compound method of claim 2 wherein the metallic radioisotope is  $Tc^{99}$ ,  $In^{111}$  or  $Gd^{153}$ .
- 4. (currently amended) The compound method of claim 1 wherein C is the residue of the (b)-monocarboxylic acid.

- 5. (currently amended) The compound method of claim 4 wherein Y is a divalent monomer, dimer or trimer of  $N(H)(CH_2)_{2-6}N(H)$ .
- 6. (currently amended) The compound method of claim 5 wherein Y is --N(H)(CH<sub>2</sub>)<sub>4</sub>NH--.
- 7. (currently amended) The <del>compound</del> <u>method</u> of claim 1 wherein Det is EDTA, DTPA, DOTA, TETA, or DCTA.
- 8. (currently amended) The compound method of claim 3 wherein Det comprises DTPA.
- 9 13 (cancelled).
- 14. (currently amended) A <u>The</u> method of <u>claim 1</u>, <u>evaluating kidney</u>, <u>liver</u>, <u>spleen or intestinal function in a mammal</u> comprising administering to said mammal a detectable amount of a <u>the</u> compound of <u>claim 1</u> in combination with a pharmaceutically acceptable vehicle, and detecting the presence of said compound in <u>the a tumor of the</u> kidney, liver, pancreas, spleen, or intestine of said mammal.
- 15. (original) The method of claim 14 wherein the administration is parenteral.
- 16. (original) The method of claim 15 wherein the administration is intravenous.
- 17. (original) The method of claim 16 wherein the administration is intraperitoneal.
- 18. (original) The method of claim 14 wherein the administration is oral.
- 19. (cancelled)

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- 20. (currently amended) The method of claim  $\frac{19}{1}$  wherein the administration is parenteral.
- 21. (currently amended) The method of claim 19 1 wherein the administration is oral.
- 22. (currently amended) The method of claim 19 14 wherein the vehicle is an aqueous vehicle.
- 23. (currently amended) The method of claim  $\frac{19}{2}$  wherein the tumor is a liver, kidney, splenic, pancreatic, or gastrointestinal tumor.
- 24. (new) The method of claim 2 wherein the metallic radioisotope is selected from the group consisting of Antimony-124, Antimony-125, Arsenic-74, Barium-103, Barium-140, Beryllium-7, Bismuth-206, Bismuth-207, Cadmium-109, Cadmium-115m, Calcium-45, Cerium-139, Cerium-141, Cerium-144, Cesium-137, Chromium-51, Cobalt-56, Cobalt-57, Cobalt-58, Cobalt-60, Cobalt-64, Erbium-169, Europium-152, Gadolinium-153, Gold-195, Gold-199, Hafnium-175, Hafnium-175-181, Indium-111, Iridium-192, Iron55, Iron-59, Krypton-85, Lead-210, Manganese-54, Mercury-197, Mercury-203, Molybdenum-99, Neodymium-147, Neptunium-237, Nickel-63, Niobiumo-95, Osmium-185+191, Palladium-103, Platinum-195m, Praseodymium-143, Promethium-147, Protactinium-233, Radium-226, Rhenium-186, Rubidium-86, Ruthenium 103, Ruthenium - 106, Scandium - 44, Scandium - 46, Selenium - 75, Silver - 110m, Silver-111, Sodium-22, Strontium-85, Strontium-89, Strontium-90, Sulfur-35, Tantalum-182, Technetium-99m, Tellurium-125, Tellurium-132, Thallium-204, Thorium-228, Thorium-232, Thallium-170, Tin-113, Titanium-44, Tungsten-185, Vanadium-48, Vanadium-49, Ytterbium-169, Yttrium-88, Yttrium-90, Yttrium-91, Zinc-65, and Zirconium-95.

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25. (New) The method of claim 1 wherein Det comprises a chelating group selected from the group consisting of EDTA, DTPA, DOTA, TETA, DCTA, 15N4, 12N3, 2-p-nitrobenzyl-1,4,7,10-tetraazacyclododecane-N,N',N",N"'-tetraacetic acid, and BAT.